

Remarks

This AMENDMENT is in response to the Official Action dated 03/06/06. A Petition for a ONE-MONTH Extension of Time is submitted to extend the time for response to 07/06/06.

Here is an overview of the status of all the claims in the present case as a result of the present AMENDMENT.

Claims 25, 26, 28, 29, 31, 38, 40, 44, 47, 49, 50, 52, and 53 are currently in this case.

Claims 1-24, 27, 30, 32-37, 39, 41-43, 45, 46, 48, and 51 are cancelled.

Claims 25, 28, 29, 31, 38, 40, 44, 47, 49, 50, and 52 are currently amended herein.

Claim 53 is newly submitted herein.

Claims 26 remains in the case as originally filed.

The claims in the case which are independent claims are claims 25, 52, and 53.

Original claims and previously amended claims are ultimately dependent upon currently amended independent Claim 25.

Now turning to objections and rejections in the Official Action dated 03/06/06, Examiner Fernandez objected to claim 28 for containing redundant language. That redundant language has been deleted herein. Therefore, it is respectfully submitted that the ground for objection to claim 28 has been removed.

In the Official Action dated 03/06/06, Examiner Fernandez rejected claim 42 under 35 USC § 102 as being anticipated by Wang (6,514,762), as not being covered by the Applicants Provisional Application 60/117,755.

In addition, Examiner Fernandez rejected claims 30 and 42 under 35 USC § 112.

Herein claims 30 and 42 are cancelled.

In the Official Action dated 03/06/06, Examiner Fernandez rejected the remainder of the Applicant's claims under 35 USC § 103, as being unpatentable over Weidlich et al (5,103,837) in view of Hofmann (6,009,347), either as a pair of references together, or in combination with other references including Zewert et al (5,749,847), Widera et al (Journal of Immunology, 2000, 164:4635-4640), or Lerner (WO 97/18855).

Among the references cited by Examiner Fernandez in the Official Action dated 03/06/06, the only reference not previously discussed extensively by the Applicant's representative is Weidlich et al (5,103,837).

In this respect, Weidlich et al (5,103,837) discloses a polymer (which is a macromolecule) which serves as a carrier for a steroid, which is a molecule (but not a macromolecule). Only the carried steroid (which is not a macromolecule) is delivered into the cells by diffusion. There is no teaching, and there is no reason why the carrier polymer, which is a macromolecule, would itself be delivered into the cells. This is in sharp

contrast with the Applicant's currently claimed invention (as claims 25, 52, and 53 are currently reciting) wherein the macromolecules themselves are delivered into the cells.

Moreover, with Weidlich et al (5,103,837), the non-macromolecular steroid is not delivered into the cells by the applied electric field. Instead, the steroid is delivered into the cells by diffusion. Moreover, the electrode is relevant only with respect to a rise in a stimulation threshold, such as for a heart pacemaker, not with respect to delivering releasable macromolecules into cells, as provided by the Applicant's currently claimed invention.

The Abstract of Weidlich et al (5,103,837) recites the use and environment of the electrode as follows, "The electrode reduces the postoperative increase in the stimulation threshold, as well as the growth of connective tissue, without any reduction in capacitance". Also, see claim 1 of Weidlich et al (5,103,837) which sets forth the essence of said patent.

Moreover, there is no reason to deliver any of the disclosed Weidlich et al (5,103,837) polymers (macromolecules) into the cells because the disclosed polymers have no therapeutic function. For example, the polymer sulfonated polytetrafluorethylene (see claim 2) has no disclosed therapeutic function.

Generally, the Weidlich et al (5,103,837) polymers are used to provide a smooth surface to prevent cellular damage to heart tissue. The purpose of the Weidlich et al (5,103,837) polymers

is diametrically opposed to delivering the polymers into heart cells. On the contrary, the Weidlich et al (5,103,837) polymers are used to minimize interaction with heart cells.

In review, clearly Weidlich et al (5,103,837) do not disclose delivering "macromolecules" into biological cells as provided by the Applicant's currently claimed invention.

Moreover Weidlich et al (5,103,837) do not disclose delivering any materials into biological cells "in the tissues penetrated by the electrode, by an applied electric field applied to the tissues penetrated by the electrode" as provided by the Applicant's currently claimed invention.

Now turning to Hofmann (6,009,347), the Hofmann patent is directed to the arrangement and spacing of needle electrodes and the voltages applied to those needle electrodes. The needle electrodes are placed in a grid, and the voltages can be applied in a rotating pattern. There are two types of needle electrodes, both of which are used to penetrate tissues. One type of needle electrode is hollow, like a hypodermic needle; and the other type of needle electrode is solid throughout. The hollow needles are used to inject a treatment agent into the tissue to be treated.

By having hollow needles used for injection of drugs into tissues, there is clearly no need for any Hofmann electrode to be pre-coated with a "coating having at least one static layer of releasable macromolecules to be delivered into biological cells" such as provided by the Applicant's currently claimed invention.

With the Hofmann (6,009,347) electrodes, having a pre-coated electrode would be unnecessary and superfluous.

In addition, there is no disclosure in Hofmann (6,009,347) of eliminating the hollow needles used for drug injection. Also, there is no disclosure in Hofmann (6,009,347) of using any kind of substitute for the hollow needles used for drug injection. With the use of hollow needles for drug injection, there is simply no motivation to have pre-coated electrodes that are coated with macromolecules for delivery into biological cells.

As stated in Hofmann (6,009,347), referring specifically to FIG. 2, the illustrated connector template is shown in use in treatment of a prostate cancer or the like. In this instance, the connector 22 is shown mounted on an elongated support rod 54 of an ultra-sound probe 56 which is shown inserted into the rectum of a patient. The sound probe is used to visualize the prostate and the location of the electrodes in the prostate. The template is then in a position such that a plurality of needle electrodes 58, 60 and 62 in a first row are inserted through three of the horizontal through bores, as illustrated, and into the prostate of the patient. In this instance, two of the needle electrodes, 58 and 62, are illustrated as being solid needle electrodes and a center electrode 60 is shown to be hollow to enable the injection of molecules, such as a drug or therapeutic agent or other material [emphasis added].

A second, or lower row of needle electrodes 64, 66 and 68 is directly below the aforementioned electrodes and extend through

the through bores of the connector template and into the prostate of the patient. In this instance, two outer needles, 64 and 68, are hollow to enable the injection of a therapeutic or other agent into the prostate of the patient. [emphasis added]. These may be left in place following the injection of the therapeutic agent and serve as the electrodes for the application of the electrical pulses to the tissue of the prostate or cancer cells within the prostate.

The hollow needles 60, 64 and 68 have outlet ports at the tip, as illustrated. For example, needle 64 is shown to have outlet ports 70 and 72. Similarly, outlet ports in needles 60 and 68 are shown but not given reference numerals. [emphasis added]

Clearly, Hofmann (6,009,347) does not contemplate any type of a pre-coated electrode for administration of any drug for electroporation thereof.

As a matter of fact, Hofmann (6,009,347) discusses the prior art of electroporation with electrodes, and nowhere in the discussion of the prior art is there a disclosure of the Applicant's currently claimed invention of electrodes being coated with a coating having at least one static layer of releasable macromolecules to be delivered into biological cells.

More specifically, in the part of the specification of Hofmann (6,009,347) which discusses the prior art, Hofmann (6,009,347) states that with in vivo applications of electroporation, electrodes are provided in various configurations such as, for example, a caliper that grips the

epidermis overlying a region of cells to be treated.

Alternatively, needle-shaped electrodes may be inserted into the patient, to access more deeply located cells. In either case, after the implant agent is injected into the treatment region, the electrodes apply an electrical field to the region. [emphasis added].

Hofmann (6,009,347) also states that a number of experiments have been conducted to test therapeutic application of electroporation for cancer treatment in a process now termed electrochemotherapy. This treatment is carried out by infusing an anticancer drug directly into the tumor and applying an electric field to the tumor between a pair of electrodes. [emphasis added].

In review, Hofmann (6,009,347) does not teach the use of a needle electrode, that is penetrated into tissue, that is pre-coated with any tissue treating agent, let alone a macromolecular treating agent. Moreover, in the four corners of Hofmann (6,009,347), there is no teaching of any way to administer a drug to be electroporated into biological cells in vivo other than the use of a hypodermic needle, or the like, to inject or infuse a treating material into a tissue to be treated. Such limited treatment means disclosed in Hofmann (6,009,347), to a person with ordinary skill in the art, clearly teach away from employing a pre-coated needle electrode to penetrate tissues for administering a macromolecule thereto.

In view of the above, the combination of Weidlich et al (5,103,837) and Hofmann (6,009,347) should not prevent the allowance of the Applicant's currently claimed invention.

Turning to Zewert et al (5,749,847), Widera et al (Journal of Immunology, 2000, 164:4635-4640), and Lerner (WO 97/18855), the following remarks are presented.

Zewert et al (5,749,847) disclose non-penetrating electrodes and a process of electroporation that is used to move a nucleotide component on the non-penetrating electrodes past the dead cells of the stratum corneum into an organism. Moreover, with Zewert et al (5,749,847), once the nucleotide component is in the organism below the stratum corneum, the nucleotide component resides in the interstitial spaces between the cells of the organism and does not penetrate into the cells. Therefore, the non-penetrating electrodes of Zewert et al (5,749,847) do not cause the nucleotide component to be delivered into the cells of the organism.

As stated above, with Zewert et al (5,749,847), the electrodes that are employed for the electroporation do not penetrate into tissues. Instead, the Zewert et al (5,749,847) electrodes simply sit on the surface of the stratum corneum. The Zewert et al (5,749,847) electrodes are not present in the tissues that underlie the stratum corneum. This is in sharp contrast with the Applicant's currently claimed invention wherein the electrode is for penetration into tissues, and releasable

macromolecules from a static layer coating on the electrode are delivered into biological cells in the penetrated tissues.

Clearly, Weidlich et al (5,103,837), Hofmann (6,009,347), or Zewert et al (5,749,847), either alone or in combination, should not be used to reject the Applicant's currently claimed invention. In this respect, it is respectfully submitted that the grounds for rejection of the Applicant's currently claimed invention based on Weidlich et al (5,103,837), Hofmann (6,009,347), or Zewert et al (5,749,847) be reconsidered and removed.

Turning to Widera et al (Journal of Immunology, 2000, 164:4635-4640), Widera et al use a three-step process for delivering a DNA vaccine into biological cells using three types of apparatuses. The first step is the use of a hypodermic needle (the first apparatus) to inject the DNA vaccine into tissues. The second step is to penetrate an electrode array (the second apparatus) into the tissues. The third step is to use an electric field generating apparatus (the third apparatus) to apply electric fields to the electrode array. The three-step process using the three types of apparatuses is substantially the same as the three-step process using three types of apparatuses that is set forth U. S. Patent No. 5,273,525 of Hofmann which is disclosed on page 3 of the Applicant's specification.

However, these teachings of Widera et al (Journal of Immunology) and U. S. Patent No. 5,273,525 of Hofmann are in

sharp contrast with the Applicant's currently claimed invention wherein only two steps are used with only two apparatuses. More specifically, the first apparatus is the Applicant's electrode that is pre-coated with macromolecules; and the second apparatus is an apparatus for generating electric fields that are applied to the pre-coated electrode having a "coating having at least one static layer of releasable macromolecules to be delivered into biological cells".

With the Applicant's currently claimed invention, the first step is to penetrate tissues with the coated electrode; and the second step is to apply electric field onto the coated electrode in the penetrated tissues.

Clearly, Weidlich et al (5,103,837), Hofmann (6,009,347), or Widera et al (Journal of Immunology, 2000, 164:4635-4640), either alone or in combination, should not be used to reject the Applicant's currently claimed invention. In this respect, it is respectfully submitted that the grounds for rejection of the Applicant's currently claimed invention based on Weidlich et al (5,103,837), Hofmann (6,009,347), or Widera et al (Journal of Immunology, 2000, 164:4635-4640) be reconsidered and removed.

Turning to Lerner (WO 97/18855), Lerner (WO 97/18855) discloses electrodes (see page 26, lines 22-38) that are designed to have smooth, non-penetrating surfaces to deliver material into tissues or organs by iontophoresis. Iontophoresis does not accomplish the critical next step of getting material into cells

using electroporation. There are many examples of iontophoresis being used for delivery of drugs into tissue. However, iontophoresis is not electroporation.

The tissues treated by Lerner (WO 97/18855) are not penetrated by the Lerner electrodes. Instead, the Lerner electrodes are placed on surfaces (such as skin) or are inserted (not penetrated) into blood vessels and body cavities where tissue treatment takes place. In addition, Lerner teaches that normal organ activity takes place while the Lerner electrode is inserted. For example, on page 29, lines 25-32, there is a teaching that a nostril-inserted electrode has a hole in it so that normal breathing can be conducted.

Clearly, Lerner does not teach the Applicant's currently claimed invention which provides "A needle electrode for penetration into tissues which includes a coating having at least one static layer of releasable macromolecules to be delivered into biological cells, in the tissues penetrated by the electrode, by an applied electric field applied to the tissues penetrated by the electrode".

Clearly, Weidlich et al (5,103,837), Hofmann (6,009,347), or Lerner (WO 97/18855), either alone or in combination, should not be used to reject the Applicant's currently claimed invention. In this respect, it is respectfully submitted that the grounds for rejection of the Applicant's currently claimed invention based on Weidlich et al (5,103,837), Hofmann (6,009,347), or Lerner (WO 97/18855) be reconsidered and removed.

No additional fees are required with respect to the type or number of claims.

A PETITION FOR REQUEST FOR EXTENSION OF TIME is filed concurrently herewith, along with a payment in the amount of \$60.00.

In view of the foregoing, it is respectfully requested that claims 25, 26, 28, 29, 31, 38, 40, 44, 47, 49, 50, 52, and 53 be deemed allowable. If the Examiner believes otherwise, or has any comments or questions, or has any suggestions for putting the case in condition for final allowance, the Examiner is respectfully urged to contact the undersigned attorney of record at the telephone number below, so that an expeditious resolution may be effected and the case passed to issue promptly.

Respectfully submitted,

July 6, 2006
Date

Marvin S. Townsend
Marvin S. Townsend
Registration Number 27,959
Attorney for Applicant

Marvin S. Townsend
Patent Attorney
8 Grovepoint Court
Rockville, MD 20854
(Voice and Fax) 301-279-0660
E-mail: MTowsend@aol.com

Certificate of Mailing

I hereby certify that this correspondence is being deposited with the United States Postal Service as First Class Mail, with sufficient postage, in an envelope addressed to:

Mail Stop NON-FEE AMENDMENT
Commissioner for Patents
P. O. Box 1450
Alexandria, VA 22313-1450

Name of person making the deposit, ;

Signature, ;

Date, ;

Date, ;